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XX 19-NOV-1998; 99US-0113635.
PR 22-DEC-1998; 98US-0113635.
PR 07-APR-1999; 99US-0128194.
XX (INCY-) INCYTE PHARM INC.
PA Yue H, Tang YT, Corley NC, Guegler KJ, Gorgone GA, Baughn MR;
XX Lu DAM, Lal P, Hillman JL, Yang J;
PI WPI; 2000-387796/33.
DR N-PSDB; AAA27386.
XX Immunoglobulin superfamily proteins, the agonist and antagonist of the
PT protein is useful for preventing and treating disorders associated with
PT altered levels of the protein such as cancer, immune system disorders
PT -
PS Claim 1; Page 82-83; 105pp; English.
XX The present sequence is the human immunoglobulin superfamily protein
CC IGFAM-6. Its gene was isolated from a cDNA library of leg
CC tissue. It is expressed in reproductive, nervous and
CC cardiovascular tissue, where cancer and inflammation are common. The
CC gene, protein, its antibodies, agonists and antagonists are suitable for
CC diagnosing and treating many diseases, including cancer, immune system
CC disorders (such as inflammation, AIDS, allergies, anaemia, Crohn's
CC arteriosclerosis, asthma, atherosclerosis, cholecystitis, Crohn's
CC disease, diabetes mellitus, emphysema, Graves' disease, hepatitis,
CC multiple sclerosis, psoriasis, rheumatoid arthritis, scleroderma,
CC systemic lupus erythematosus and ulcerative colitis), complications of
CC cancer, haemodialysis and extracorporeal circulation, trauma and
CC haematopoietic cancer (such as leukaemia) and infections caused by
CC bacteria, viruses, fungi or parasites.
XX Sequence 310 AA;
SQ Query Match 100.0%; Score 1635; DB 21; Length 310;
Best Local Similarity 100.0%; Pred. No. 2.2e-133; Indels 0; Gaps 0;
Matches 310; Conservative 0; Mismatches 0;
QY 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQEVEVDL 240
QY 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
QY 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310
RESULT 2
AAB27276
ID AAB27276 standard; Protein; 310 AA.
XX AAB27276;
XX 23-FEB-2001 (first entry)
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XX Human confluency regulated adhesion molecule 1 #2.
DE Immunoglobulin superfamily; Ig Sf; vascular adhesion molecule;
XX inflammation; cancer; wound; angiogenesis; human;
KW confluency regulated adhesion molecule 1; CRAM-1; JAM-2.
XX Homo sapiens.
OS WO200053749-A2.
PN 14-SEP-2000.
PD 13-MAR-2000; 2000WO-EP02219.
XX 11-MAR-1999; 99EP-0200746.
PR (RMFD-) RMF DICTAGENE SA.
XX Imhof BA, Aurrand-Lions M;
PI WPI; 2000-587436/55.
XX N-PSDB; AAA95306.
DR Isolated human Confluency Regulated Adhesion Molecule 1 or 2 (CRAM-1 or
PT CRAM-2) polypeptide, useful for treatment of tumors, inflammation
PT reactions and modulating vascular permeability -
XX Claim 2; Fig 6; 59pp; English.
PS The present sequence is the human confluency regulated adhesion molecule
XX 1 (CRAM-1, also known as JAM-2). CRAM-1 is one of the vascular adhesion
CC proteins of the immunoglobulin superfamily (Ig Sf). The CRAM-1 protein
CC and coding sequence can be used in the treatment of cancer, inflammation,
CC to modulate cell-cell interactions and angiogenesis, and in the
CC modulation of wound healing.
XX Sequence 310 AA;
SQ Query Match 99.6%; Score 1629; DB 21; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQEVEVDL 240
QY 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
QY 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310
RESULT 3
AAB33457
ID AAB33457 standard; Protein; 310 AA.
XX AAB33457;
XX AC AAB33457;
```

XX DT 29-JAN-2001 (first entry)

XX DE Human PRO1868 protein UNQ859 SEQ ID NO:193.

XX KW Human; immune related disease; diagnosis; antiinflammatory; cardiant;

XX KW haematological; antiarthritic; antirheumatic; immunosuppressive;

XX KW haemostatic; antithyroid; antidiabetic; nootropic; neuroprotective;

XX KW antianemic; hepatotropic; virucide; antipsoriatic; antiallergic;

XX KW antiasthmatic; systemic lupus erythematosus; rheumatoid arthritis;

XX KW osteoarthritis; spondyloarthropathy; systemic sclerosis; sarcoidosis;

XX KW idiopathic inflammatory myopathy; Sjogren's syndrome; thyroiditis;

XX KW systemic vasculitis; autoimmune haemolytic anaemia; diabetes mellitus;

XX KW autoimmune thrombocytopenia; immune-mediated renal disease;

XX KW demyelinating disease; hepatobiliary disease; Whipple's disease;

XX KW inflammatory bowel disease; gluten-sensitive enteropathy;

XX KW autoimmune disease; immune-mediated skin disease; allergic disease;

XX KW immunological disease; transplantation associated disease;

XX KW graft rejection; graft-versus-host-disease.

OS Homo sapiens.

XX KW WO200053759-A2.

PN 14-SEP-2000.

XX 02-MAR-2000; 2000WO-US05841.

PR 08-MAR-1999; 99WO-US05028.

PR 10-MAR-1999; 99US-0123618.

PR 12-MAR-1999; 99US-0123957.

PR 23-MAR-1999; 99US-0125775.

PR 12-APR-1999; 99US-0128849.

PR 20-APR-1999; 99WO-US08615.

PR 28-APR-1999; 99US-0134445.

PR 04-MAY-1999; 99US-0132371.

PR 14-MAY-1999; 99US-0134287.

PR 02-JUN-1999; 99WO-US12252.

PR 23-JUN-1999; 99US-0141037.

PR 26-JUL-1999; 99US-0144758.

PR 26-JUL-1999; 99US-0145698.

PR 28-JUL-1999; 99US-0146222.

PR 01-SEP-1999; 99WO-US20111.

PR 08-SEP-1999; 99WO-US20594.

PR 13-SEP-1999; 99WO-US20944.

PR 15-SEP-1999; 99WO-US21090.

PR 15-SEP-1999; 99WO-US21547.

PR 05-OCT-1999; 99WO-US23089.

PR 29-OCT-1999; 99US-0162506.

PR 29-NOV-1999; 99WO-US28214.

PR 30-NOV-1999; 99WO-US28313.

PR 30-NOV-1999; 99WO-US28409.

PR 01-DEC-1999; 99WO-US28301.

PR 01-DEC-1999; 99WO-US28634.

PR 02-DEC-1999; 99WO-US28551.

PR 02-DEC-1999; 99WO-US28564.

PR 02-DEC-1999; 99WO-US28565.

PR 16-DEC-1999; 99WO-US30095.

PR 20-DEC-1999; 99WO-US30999.

PR 30-DEC-1999; 99WO-US31274.

PR 05-JAN-2000; 2000WO-US00219.

PR 06-JAN-2000; 2000WO-US00277.

PR 06-JAN-2000; 2000WO-US00376.

PR 11-FEB-2000; 2000WO-US03565.

PR 18-FEB-2000; 2000WO-US04341.

PR 18-FEB-2000; 2000WO-US04342.

PR 22-FEB-2000; 2000WO-US04414.

XX (GETH) GENENTECH INC.

XX Ashkenazi AJ, Baker KP, Goddard A, Gurney AL, Hebert C, Henzel W;

PI Kabakoff RC, Lu Y, Pan J, Pennica D, Shelton DL, Smith V;

PI Stewart TA, Tumas D, Watanabe CK, Wood WI, Yan M;

XX WPI; 2000-572271/53.

DR N-PSDB; AAC58622.

XX

PT Sixty four PRO polypeptides, useful in the diagnosis and treatment of

PT immune related disorders, e.g. systemic lupus erythematosus, rheumatoid

PT arthritis, osteoarthritis, thyroiditis and diabetes mellitus -

PS Claim 33; Fig 88; 309pp; English.

XX

CC The present invention describes sixty four human PRO proteins which can

CC be used in the treatment of immune related diseases. The human PRO

CC proteins, anti-PRO antibodies, agonists and antagonists are useful for

CC treating and diagnosing immune related disorders. The disorders are

CC selected from systemic lupus erythematosus, rheumatoid arthritis,

CC osteoarthritis, juvenile chronic arthritis, spondyloarthropathies,

CC systemic sclerosis, idiopathic inflammatory myopathies, Sjogren's

CC syndrome, systemic vasculitis, sarcoidosis, autoimmune haemolytic

CC anaemia, autoimmune thrombocytopenia, thyroiditis, diabetes mellitus,

CC immune-mediated renal disease, demyelinating diseases of the central

CC and peripheral nervous systems, hepatobiliary diseases, inflammatory

CC bowel disease, gluten-sensitive enteropathy and Whipple's disease,

CC autoimmune or immune-mediated skin diseases, allergic diseases,

CC immunological diseases of the lung, and transplantation associated

CC diseases including graft rejection and graft-versus-host-disease.

CC AAC58397 to AAC58578 represent PCR primers and hybridisation probes used

CC in the isolation of human PRO sequences. AAC58579 to AAC58642 and

CC AAB33414 to AAB33477 represent human PRO polynucleotide and protein

CC sequences given in the exemplification of the present invention.

XX

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 21; Length 310;

Best Local Similarity 99.7%; Pred. No. 7.2e-133; Indels 0; Gaps 0;

Matches 309; Conservative 0; Mismatches 1;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60

DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60

QY 61 SDPRIWKKIODEQTTVFFDNKIQGLAGRAEILGKTSKIWNVTRRDSALYRCEVVAR 120

DB 61 SDPRIWKKIODEQTTVFFDNKIQGLAGRAEILGKTSKIWNVTRRDSALYRCEVVAR 120

QY 121 NDRKEIDEIVIELTVQVKPVPVCRKAVPVGKMATLHCQESGHPHYSWYRNDVPL 180

DB 121 NDRKEIDEIVIELTVQVKPVPVCRKAVPVGKMATLHCQESGHPHYSWYRNDVPL 180

QY 181 PTDSRANPRFRNSSSHLNSETGLVFTAVHKDDSGQYVCIASNDAGSARCEQEMEVYDL 240

DB 181 PTDSRANPRFRNSSSHLNSETGLVFTAVHKDDSGQYVCIASNDAGSARCEQEMEVYDL 240

QY 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQGESYKNPKGPDGWNVYRTDEEG 300

DB 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQGESYKNPKGPDGWNVYRTDEEG 300

QY 301 DFRHKSSRFVI 310

DB 301 DFRHKSSRFVI 310

RESULT 4

AA96735

ID AA96735 standard; Protein; 310 AA.

XX

AC AA96735;

XX

DT 26-SEP-2000 (first entry)

XX

DE PRO1868, an A33 antigen homologue.

XX

KW PRO1868; A33 antigen; secreted protein; transmembrane protein;

KW anti-inflammatory; cytostatic; recombinant production; gene therapy.

XX OS Homo sapiens.
XX FH Key Location/Qualifiers
XX FT Peptide 1..30
XX FT /label= Signal_peptide
XX FT Modified-site 26..31
XX FT /note= "N-myristoylation site"
XX FT Modified-site 69..77
XX FT /note= "Tyrosine kinase phosphorylation site"
XX FT Modified-site 104..107
XX FT /note= "N-glycosylation site"
XX FT Modified-site 106..109
XX FT /note= "Casein kinase II phosphorylation site"
XX FT Modified-site 107..110
XX FT /note= "cAMP- and cGMP-dependent protein kinase phosphorylation site"
XX FT Modified-site 192..195
XX FT /note= "N-glycosylation site"
XX FT Modified-site 215..220
XX FT /note= "N-myristoylation site"
XX FT Modified-site 226..231
XX FT /note= "N-myristoylation site"
XX FT Domain 243..263
XX FT /label= Transmembrane_domain
XX FT Modified-site 243..248
XX FT /note= "N-myristoylation site"
XX FT Modified-site 244..249
XX FT /note= "N-myristoylation site"
XX FT Modified-site 262..267
XX FT /note= "N-myristoylation site"
XX FT Modified-site 296..299
XX FT /note= "Casein kinase II phosphorylation site"
XX PN WO200036102-A2.
XX PD 22-JUN-2000.
XX PF 01-DEC-1999; 99WO-US28634.
XX PR 16-DEC-1998; 98US-0112851.
XX PR 16-DEC-1998; 98US-0113145.
XX PR 22-DEC-1998; 98US-0113511.
XX PR 12-JAN-1999; 99US-0115558.
XX PR 12-JAN-1999; 99US-0115565.
XX PR 12-JAN-1999; 99US-0115733.
XX PR 09-FEB-1999; 99US-0119341.
XX PR 10-FEB-1999; 99US-0119537.
XX PR 12-FEB-1999; 99US-0119965.
XX PR 02-JUN-1999; 99WO-US12252.
XX PA (GETH) GENENTECH INC.
XX PI Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
XX PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
XX PI Wood WI;
XX XX
XX WPI; 2000-431586/37.
XX DR N-PSDB; AAA51265.
XX XX
XX Isolated nucleic acid molecule encodes a PRO polypeptide which is a
XX transmembrane polypeptide
XX Claim 1; Fig 14; 154pp; English.
XX This is PRO1868, a putative homologue of A33 antigen, a known
XX colorectal cancer-associated marker. The invention concerns novel
XX secreted and transmembrane proteins, designated PRO polypeptides. The
XX cDNA and gene sequences are useful in the recombinant production of PRO
XX polypeptides, as a hybridization probe to screen libraries to isolate
XX cDNAs with sequence identity to PRO polypeptides or to map the gene
XX encoding the PRO polypeptides and analyzing genetic disorders. The
XX cDNA/gene can also be used to produce transgenic animals useful for the

CC development and screening of therapeutically useful reagents. They can
CC also be used in gene therapy, e.g. to replace a defective gene.
XX SQ Sequence 310 AA;
Query Match 99.6%; Score 1629; DB 21; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVVQEFSEVLSCLITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVVQEFSEVLSCLITDSQT 60
Qy 61 SDPRIWKIQDEQTTTVVFFDNKIQGLAGRAEILGKTSKIMNVTTRDSALYRCEVVAR 120
Db 61 SDPRIWKIQDEQTTTVVFFDNKIQGLAGRAEILGKTSKIMNVTTRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVKAIVPGVGMATLHCQESGHPHPRHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVKAIVPGVGMATLHCQESGHPHPRHYSWYRNDVPL 180
Qy 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYVCIASNDAGSARCEQEMEYVDL 240
Db 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYVCIASNDAGSARCEQEMEYVDL 240
Qy 241 NIGGIIGVLVLAVALILTLGICCAVRRGVFINNKDGSYKPKPGDGVNYIRTDEEG 300
Db 241 NIGGIIGVLVLAVALILTLGICCAVRRGVFINNKDGSYKPKPGDGVNYIRTDEEG 300
Qy 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310
RESULT 5
AAM93323
ID AAM93323 standard; Protein; 310 AA.
XX AC AAM93323;
XX DT 06-NOV-2001 (first entry)
XX DE Human polypeptide, SEQ ID NO: 2845.
XX KW Human; full length cDNA; cDNA synthesis; oligo-capping.
XX OS Homo sapiens.
XX PN EPI130094-A2.
XX PD 05-SEP-2001.
XX PF 07-JUL-2000; 2000EP-0114089.
XX PR 08-JUL-1999; 99JP-0194486.
XX PR 11-JAN-2000; 2000JP-0118774.
XX PR 02-MAY-2000; 2000JP-0193765.
XX PA (HELI-) HELIX RES INST.
XX PI Ota T, Nishikawa T, Isogai T, Hayashi K, Ishii S, Kawai Y;
XX PI Wakamatsu A, Sugiyama T, Nagai K, Kojima S, Otsuki T, Koga H;
XX DR WPI; 2001-524255/58.
XX DR N-PSDB; AAK94243.
XX PT 830 Primers useful for synthesizing full length cDNA clones and their
XX use in genetic manipulation -
XX PS Claim 8; SEQ ID NO 2845; 1380pp + sequence listing; English.
XX The invention relates to primers for synthesizing full length cDNA
XX clones. 830 cDNA molecules encoding a human protein have been

CC isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA
CC molecules have been determined. Primers for synthesizing the full length
CC cDNA are useful for clarifying the function of the protein encoded by
CC the cDNA. The full length clones were obtained by construction of full
CC length enriched cDNA libraries that were synthesised by the oligo-capping
CC method. The primers enable the production of the full length cDNA easily
CC without any special methods. The present sequence is a polypeptide
CC encoded by a full length human cDNA of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in CD-ROM format directly from EPO.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Qy 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Db 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Qy 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGSYKPNPKGPDGVNVRTDEEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGSYKPNPKGPDGVNVRTDEEG 300
Qy 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310

RESULT 6
AAM93905
ID AAM93905 standard; Protein; 310 AA.

XX AC AAM93905;

XX DT 06-NOV-2001 (first entry)

XX DE Human polypeptide, SEQ ID NO: 4051.

XX KW Human; full length cDNA; cDNA synthesis; oligo-capping.

XX OS Homo sapiens.

XX PN EP1130094-A2.

XX PD 05-SEP-2001.

XX PF 07-JUL-2000; 2000EP-0114089.

XX PR 08-JUL-1999; 99JP-0194486.

XX PR 11-JAN-2000; 2000JP-0119774.

XX PR 02-MAY-2000; 2000JP-0183765.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Nishikawa T, Isogai T, Hayashi K, Ishii S, Kawai Y;

XX PI Wakamatsu A, Sugiyama T, Nagai K, Kojima S, Otsuki T, Koga H;

XX WIPI; 2001-524255/58.

XX DR N-PSDB; AAK94867.

XX 830 Primers useful for synthesizing full length cDNA clones and their
PT use in genetic manipulation -
XX
XX Claim 8; SEQ ID NO 4051; 1380pp + sequence listing; English.
XX
CC The invention relates to primers for synthesising full length cDNA
CC clones. 830 cDNA molecules encoding a human protein have been
CC isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA
CC molecules have been determined. Primers for synthesising the full length
CC cDNA are useful for clarifying the function of the protein encoded by
CC the cDNA. The full length clones were obtained by construction of full
CC length enriched cDNA libraries that were synthesised by the oligo-capping
CC method. The primers enable the production of the full length cDNA easily
CC without any special methods. The present sequence is a polypeptide
CC encoded by a full length human cDNA of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in CD-ROM format directly from EPO.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Qy 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Db 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Qy 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGSYKPNPKGPDGVNVRTDEEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGSYKPNPKGPDGVNVRTDEEG 300
Qy 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310

RESULT 7

AAM12440

ID AAM12440 standard; Protein; 310 AA.

XX AC AAM12440;

XX DT 24-OCT-2001 (first entry)

XX DE Human PRO1868 polypeptide sequence.

XX KW Human secretory and transmembrane; PRO; mammalian; cancer; lung;
KW breast; prostate; cervical; tumour necrosis factor-alpha; TNF-alpha;
KW cartilage; ear; proliferation; glucose; free fatty acid; skeletal muscle;
KW adipocyte; A-peptide; factor VIIa; gene therapy.

XX OS Homo sapiens.

XX PN WO200140466-A2.

XX PD 07-JUN-2001.

XX PF 01-DEC-2000; 2000WO-US32678.

XX

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PR 01-DEC-1999; 99WO-US28301.
PR 01-DEC-1999; 99WO-US28634.
PR 02-DEC-1999; 99WO-US28551.
PR 02-DEC-1999; 99WO-US28564.
PR 02-DEC-1999; 99WO-US28565.
PR 09-DEC-1999; 99WO-US30095.
PR 16-DEC-1999; 99WO-US30911.
PR 20-DEC-1999; 99WO-US30999.
PR 30-DEC-1999; 99WO-US31243.
PR 06-JAN-2000; 2000WO-US00277.
PR 06-JAN-2000; 2000WO-US00376.
PR 11-FEB-2000; 2000WO-US03565.
PR 18-FEB-2000; 2000WO-US04341.
PR 18-FEB-2000; 2000WO-US04342.
PR 22-FEB-2000; 2000WO-US04414.
PR 24-FEB-2000; 2000WO-US04914.
PR 24-FEB-2000; 2000WO-US05004.
PR 01-MAR-2000; 2000WO-US07377.
PR 20-MAR-2000; 2000WO-US07377.
PR 21-MAR-2000; 2000WO-US07532.
PR 30-MAR-2000; 2000WO-US08439.
PR 17-MAY-2000; 2000WO-US13705.
PR 22-MAY-2000; 2000WO-US14042.
PR 30-MAY-2000; 2000WO-US14941.
PR 02-JUN-2000; 2000WO-US15264.
PR 10-NOV-2000; 2000WO-US30873.
XX
PA (GETH ) GENENTECH INC.
XX
XX Baker KP, Beresini M, DeForge L, Desnoyers L, Filvaroff E, Gao W;
PI Gerritsen ME, Goddard A, Godowski PJ, Gurney AL, Sherwood S;
PI Smith V, Stewart TA, Tumas D, Watanabe CK, Wood WI, Zhang Z;
XX
XX WPI; 2001-408281/43.
XX N-PSDB; AAS21512.
XX
XX Isolated, secretory and transmembrane PRO polypeptide used to detect
XX other PRO polypeptides, link bioactive molecules to cells expressing
XX PRO polypeptides, and detect the presence of mammalian tumours e.g.
XX lung, breast, prostate, cervical
XX
XX Claim 12; Fig 538; 813pp; English.
XX
XX AAU12172-AAU12446 represent novel human secretory and transmembrane
XX PRO polypeptides. The PRO polypeptides are useful to detect other
XX PRO polypeptides, to link bioactive molecules to cells expressing
XX PRO polypeptides, to modulate biological activities of cells expressing
XX PRO polypeptides, and to detect the presence of mammalian lung, colon,
XX breast, prostate, rectal, cervical or liver tumours by comparing PRO
XX polypeptide expression in a cell sample to that in a control sample.
XX Some of the 275 sequences are also useful to stimulate the release of
XX tumour necrosis factor-alpha (TNF-alpha) from human blood, the
XX proliferation or differentiation of chondrocytes, the proliferation or
XX gene expression in pericyte cells, the release of proteoglycans from
XX cartilage, the proliferation of inner ear utricular supporting cells or
XX of T-lymphocytes, the release of a cytokine from peripheral blood
XX monocytes (PBMCs), or the proliferation of endothelial cells. Some of
XX the PRO polypeptides may modulate glucose or free fatty acid uptake by
XX skeletal muscle cells or by adipocytes; or inhibit binding of A-peptide
XX to factor VIIA. The PRO polypeptides can be used in assays to identify
XX molecules involved in binding interactions. The polynucleotides encoding
XX PRO polypeptides can be used to generate probes, antisense RNA/DNA,
XX transgenic or knock out animals and can be used in gene therapy.
XX
XX Sequence 310 AA;
SQ
Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 MALRRPRLRLCARLPDFFLLLRGCLIGAVNLKSSNRTPVVQEFESVLSCTITDSQT 60
|||||

```

```
XX PT Sixty one nucleic acids encoding PRO polypeptides which are useful in
PT the treatment of skin diseases (e.g. psoriasis), cancers (e.g. lung
PT squamous cell carcinoma) and neurodegenerative diseases (e.g.
PT Alzheimer's disease) -
XX PS Claim 1; Fig 124; 393pp; English.
XX CC The present sequence is one of sixty one novel secreted and
CC transmembrane PRO polypeptides. The PRO polypeptides are
CC useful for treating skin diseases (e.g. psoriasis), cancers (e.g. lung
CC squamous cell carcinoma), gastrointestinal disorders (e.g.
CC enterocolitis), neurodegenerative diseases (e.g. Alzheimer's disease,
CC Parkinson's disease), wound repair, cardiovascular disorders (e.g.
CC endometrial bleeding angiogenesis, ischaemia such as coronary
CC ischaemia, atherosclerosis), inflammatory disorders (e.g. asthma,
CC rheumatoid arthritis, multiple sclerosis), infertility, AIDS and
CC diabetes and retinal disorders such as retinitis pigmentosum.
CC The PRO nucleic acids have applications in molecular biology, including
CC use as hybridization probes, and in chromosome and gene mapping.
XX SQ Sequence 310 AA;
Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFSEVLSLTIIDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFSEVLSLTIIDSQT 60
QY 61 SDPRIWKKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
DB 61 SDPRIWKKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVLTVOVKPVPVCRKAVPVGKMATLHCOESEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVLTVOVKPVPVCRKAVPVGKMATLHCOESEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEMEYVDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEMEYVDL 240
QY 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKNPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKNPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310
RESULT 9
AAB80383
ID AAB80383 standard; protein; 310 AA.
XX AC AAB80383;
XX DT 24-APR-2001 (first entry)
XX DE Secreted protein encoded by gene #13.
XX KW Secreted protein; human; autoimmune; hyperproliferation;
XX KW cardiovascular; cerebrovascular; infection; food.
XX OS Homo sapiens.
XX PN WO200107459-A1.
XX PD 01-FEB-2001.
XX PF 20-JUL-2000; 2000WO-US19735.
XX XX

PR 23-JUL-1999; 99US-0145220.
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX PI Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
XX PI Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX DR WPI; 2001-123261/13.
XX CC New isolated nucleic acid encoding 29 secreted proteins, for
XX PT diagnosing, preventing and treating e.g. autoimmune,
XX PT hyperproliferation, cardiovascular, and ocular diseases or disorders
XX PS and microorganism infections -
XX CC Claim 11; Page 538-539; 601pp; English.
XX CC The present invention relates to 29 human secreted proteins. The
XX CC invention is used to prevent autoimmune diseases e.g. rheumatoid
XX CC arthritis, hyperproliferative disorders e.g. neoplasms of the
XX CC breast or liver, cardiovascular disorders e.g. cardiac arrest,
XX CC cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
XX CC nervous system disorders e.g. Alzheimer's disease, infections
XX CC caused by bacteria, viruses and fungi and ocular disorders e.g.
XX CC corneal infection. Also used in food preparations.
XX SQ Sequence 310 AA;
Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFSEVLSLTIIDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFSEVLSLTIIDSQT 60
QY 61 SDPRIWKKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
DB 61 SDPRIWKKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVLTVOVKPVPVCRKAVPVGKMATLHCOESEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVLTVOVKPVPVCRKAVPVGKMATLHCOESEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEMEYVDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEMEYVDL 240
QY 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKNPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKNPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310
RESULT 10
AAB80408
ID AAB80408 standard; protein; 310 AA.
XX AC AAB80408;
XX DT 24-APR-2001 (first entry)
XX DE Secreted protein encoded by gene #38.
XX KW Secreted protein; human; autoimmune; hyperproliferation;
XX KW cardiovascular; cerebrovascular; infection; food.
XX OS Homo sapiens.
XX PN WO200107459-A1.
XX PF 20-JUL-2000; 2000WO-US19735.
XX XX
```

```
PD 01-FEB-2001.
XX
XX 20-JUL-2000; 2000WO-US19735.
XX
XX 23-JUL-1999; 99US-0145220.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
XX Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX WPI; 2001-123261/13.
XX
XX New isolated nucleic acid encoding 29 secreted proteins, for
XX diagnosing, preventing and treating e.g. autoimmune,
XX hyperproliferative, cardiovascular, and ocular diseases or disorders
XX and microorganism infections -
XX
XX Claim 11; Page 557-558; 601pp; English.
XX
XX The present invention relates to 29 human secreted proteins. The
XX invention is used to prevent autoimmune diseases e.g. rheumatoid
XX arthritis, hyperproliferative disorders e.g. neoplasms of the
XX breast or liver, cardiovascular disorders e.g. cardiac arrest,
XX cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
XX nervous system disorders e.g. Alzheimer's disease, infections
XX caused by bacteria, viruses and fungi and ocular disorders e.g.
XX corneal infection. Also used in food preparations.
XX
XX Sequence 310 AA;
XX
XX Query Match 99.6%; Score 1629; DB 22; Length 310;
XX Best Local Similarity 99.7%; Pred. No. 7.2e-133;
XX Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVIELTVQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
QY 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQDGESYKPNPKGPDGVNIRTDEEG 300
Db 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQDGESYKPNPKGPDGVNIRTDEEG 300
QY 301 DFRHKSSFFVI 310
Db 301 DFRHKSSFFVI 310
RESULT 11
AAB80409
ID AAB80409 standard; protein; 310 AA.
XX
XX AAB80409;
XX
XX 24-APR-2001 (first entry)
XX
XX Secreted protein encoded by gene #39.
XX
XX Secreted protein; human; autoimmune; hyperproliferation;
XX cardiovascular; cerebrovascular; infection; food.
XX
```

```
OS Homo sapiens.
XX
XX WO200107459-A1.
XX
XX 01-FEB-2001.
XX
XX 20-JUL-2000; 2000WO-US19735.
XX
XX 23-JUL-1999; 99US-0145220.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
XX Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX WPI; 2001-123261/13.
XX
XX New isolated nucleic acid encoding 29 secreted proteins, for
XX diagnosing, preventing and treating e.g. autoimmune,
XX hyperproliferative, cardiovascular, and ocular diseases or disorders
XX and microorganism infections -
XX
XX Claim 11; Page 559-560; 601pp; English.
XX
XX The present invention relates to 29 human secreted proteins. The
XX invention is used to prevent autoimmune diseases e.g. rheumatoid
XX arthritis, hyperproliferative disorders e.g. neoplasms of the
XX breast or liver, cardiovascular disorders e.g. cardiac arrest,
XX cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
XX nervous system disorders e.g. Alzheimer's disease, infections
XX caused by bacteria, viruses and fungi and ocular disorders e.g.
XX corneal infection. Also used in food preparations.
XX
XX Sequence 310 AA;
XX
XX Query Match 99.6%; Score 1629; DB 22; Length 310;
XX Best Local Similarity 99.7%; Pred. No. 7.2e-133;
XX Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVIELTVQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
QY 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQDGESYKPNPKGPDGVNIRTDEEG 300
Db 241 NIGGIIGVLVLAVALITLIGICAYRRGYFINNKQDGESYKPNPKGPDGVNIRTDEEG 300
QY 301 DFRHKSSFFVI 310
Db 301 DFRHKSSFFVI 310
RESULT 12
AAB80409
ID AAB80409 standard; protein; 310 AA.
XX
XX AAB80409;
XX
XX 29-NOV-2002 (first entry)
XX
XX Human PRO1868 polypeptide.
XX
```


XX Human; secreted and transmembrane polypeptide; PRO polypeptide;
 KW T-lymphocyte proliferation; inflammatory disease; rheumatoid arthritis;
 KW inflammatory bowel disease; Sjogren's syndrome; thyroiditis;
 KW autoimmune haemolytic anaemia; diabetes mellitus; multiple sclerosis;
 KW hepatitis; contact dermatitis; allergic disease; psoriasis; virucide;
 KW immune related disease; kidney disease; antinflammatory; antithyroid;
 KW antirheumatic; antiarthritic; immunosuppressive; antianaemic;
 KW antidiabetic; neuroprotective; hepatotropic; antiinflammatory;
 KW dermatological; antiallergic; antipsoriatic; PRO1868.
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Peptide 1..30
 FT /label= Signal_peptide
 FT Modified-site 26..31
 FT /note= "N-myristoylation site"
 FT Protein 31..310
 FT /label= Mature_PRO1868
 FT Modified-site 69..77
 FT /note= "Tyrosine kinase phosphorylation site"
 FT Modified-site 104..107
 FT /note= "N-glycosylation site"
 FT Modified-site 106..109
 FT /note= "Casein kinase II phosphorylation site"
 FT Modified-site 107..110
 FT /note= "cAMP- and cGMP-dependent protein kinase phosphorylation site"
 FT Modified-site 192..195
 FT /note= "N-glycosylation site"
 FT Modified-site 215..220
 FT /note= "N-myristoylation site"
 FT Modified-site 226..231
 FT /note= "N-myristoylation site"
 FT Domain 243..263
 FT /label= Transmembrane_domain
 FT Modified-site 243..248
 FT /note= "N-myristoylation site"
 FT Modified-site 244..249
 FT /note= "N-myristoylation site"
 FT Modified-site 262..267
 FT /note= "N-myristoylation site"
 FT Modified-site 296..299
 FT /note= "Casein kinase II phosphorylation site"
 XX
 US2002098507-A1.
 XX
 PD 25-JUL-2002.
 XX
 XX 27-DEC-2001; 2001US-0033326.
 XX
 PR 02-JUN-1999; 99WO-US12252.
 PR 01-DEC-1999; 99WO-US28634.
 PR 02-DEC-1999; 99WO-US28551.
 PR 11-FEB-2000; 2000WO-US03565.
 PR 22-FEB-2000; 2000WO-US04414.
 PR 02-MAR-2000; 2000WO-US05841.
 PR 30-MAR-2000; 2000WO-US08439.
 PR 30-MAY-2000; 2000WO-US14941.
 PR 02-JUN-2000; 2000WO-US15264.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 16-DEC-1998; 98US-113145P.
 PR 22-DEC-1998; 98US-113511P.
 PR 12-JAN-1999; 99US-115558P.
 PR 12-JAN-1999; 99US-115565P.
 PR 09-FEB-1999; 99US-115733P.
 PR 10-FEB-1999; 99US-119341P.
 PR 12-FEB-1999; 99US-119537P.
 PR 29-OCT-1999; 99US-119965P.
 PR 29-OCT-1999; 99US-162506P.
 XX
 (GETH) GENENTECH INC. PA

XX Botstein D, Deenoyers L, Ferrara N, Fong S, Gao W, Goddard A;
 PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
 PI Wood WI;
 XX WPI; 2002-673923/72.
 DR N-PSDB; ABS53477.
 XX Novel PRO polypeptides and nucleic acids encoding the polypeptides,
 PT useful for preparing a medicament for the treatment of inflammatory and
 PT immune related disorders -
 XX
 PS Claim 12; Fig 14; 125pp; English.
 XX
 CC The present invention relates to the isolation of novel human
 CC secreted and transmembrane polypeptides, designated PRO polypeptides,
 CC and the polynucleotide sequences encoding them. The PRO polypeptides
 CC of the invention include PRO1800, PRO539, PRO982, PRO1434, PRO1863,
 CC PRO1917, PRO1868, PRO3434 and PRO1927. The PRO polypeptides can
 CC inhibit the stimulation of T-lymphocyte proliferation. The PRO
 CC polypeptides are useful for the diagnosis and treatment of inflammatory
 CC diseases (e.g. inflammatory bowel disease, rheumatoid arthritis,
 CC Sjogren's syndrome, autoimmune haemolytic anaemia, thyroiditis, diabetes
 CC mellitus, multiple sclerosis, hepatitis, contact dermatitis, allergic
 CC diseases and psoriasis), immune related diseases, and kidney diseases
 CC in humans. The present sequence represents human PRO1868 polypeptide.
 XX
 SQ Sequence 310 AA;
 Query Match 99.6%; Score 1629; DB 23; Length 310;
 Best Local Similarity 99.7%; Pred. No. 7.2e-133;
 Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 MALRRPRLRLCARLPDPFELLIIFRGCLIGAVNLKSSNRTPVQVEFSEVLSCLITDSQT 60
 Db 1 MALRRPRLRLCARLPDPFELLIIFRGCLIGAVNLKSSNRTPVQVEFSEVLSCLITDSQT 60
 Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
 Db 61 SDPRIEWKKIQDEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
 Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVKAPVGVQKATLHCQSEGHPRPHYSWYRNDVPL 180
 Db 121 NDRKEIDEIVIELTVQVKPVPVCRVKAPVGVQKATLHCQSEGHPRPHYSWYRNDVPL 180
 Qy 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYYCIASNDAGSARCEEQMEVYDL 240
 Db 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYYCIASNDAGSARCEEQMEVYDL 240
 Qy 241 NIGGIIGVLVLAVALIITLIGICCAVRRGYFINNKQDGSYKPNKPGPDGVNYIRTDDEG 300
 Db 241 NIGGIIGVLVLAVALIITLIGICCAVRRGYFINNKQDGSYKPNKPGPDGVNYIRTDDEG 300
 Qy 301 DFRHKSSFVI 310
 Db 301 DFRHKSSFVI 310
 RESULT 13
 ABG91361
 ID ABG91361 standard; Protein; 310 AA.
 XX
 AC ABG91361;
 XX
 DT 29-NOV-2002 (first entry)
 XX
 DE Novel human secreted protein #7.
 XX
 KW Human; secreted protein; transmembrane protein; gene mapping;
 KW transgenic; immunogenic.
 XX
 OS Homo sapiens.
 XX

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PN US2002098505-A1.
XX 25-JUL-2002.
XX 28-DEC-2001; 2001US-0033246.
XX 02-JUN-1999; 99WO-US12252.
PR 01-DEC-1999; 99WO-US28634.
PR 02-DEC-1999; 99WO-US28551.
PR 11-FEB-2000; 2000WO-US03565.
PR 22-FEB-2000; 2000WO-US04414.
PR 02-MAR-2000; 2000WO-US05841.
PR 30-MAR-2000; 2000WO-US08439.
PR 30-MAY-2000; 2000WO-US14941.
PR 02-JUN-2000; 2000WO-US15264.
PR 01-DEC-2000; 2000WO-US32678.
PR 16-DEC-1998; 98US-113145P.
PR 22-DEC-1998; 98US-113511P.
PR 12-JAN-1999; 99US-115558P.
PR 12-JAN-1999; 99US-115565P.
PR 12-JAN-1999; 99US-115733P.
PR 09-FEB-1999; 99US-119341P.
PR 10-FEB-1999; 99US-119537P.
PR 12-FEB-1999; 99US-119565P.
PR 29-OCT-1999; 99US-162506P.
XX
PA (GETH ) GENENTECH INC.
XX
XX Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
PI Wood WI;
XX
XX WPI; 2002-665999/71.
DR N-PSDB; ABS67460.
XX
XX New human secreted and transmembrane (PRO) polypeptides, useful for
PT treating conditions requiring PRO polypeptides, for screening PRO
PT antagonists and agonists useful as drug candidates -
XX
XX Claim 12; Figure 14; 125pp; English.
XX
XX The invention relates to new human secreted and transmembrane proteins
CC (PRO) and nucleic acids of the invention. The polypeptides can be
CC administered therapeutically, especially by expressing encoding
CC polynucleotides, e.g. in therapeutic compositions. They can be used to
CC screen for PRO polypeptide antagonists and agonists useful to identify
CC drug candidates. They can also be used to produce antibodies, useful to
CC detect PRO polypeptides (e.g. diagnostically), purify PRO polypeptides or
CC therapeutically (e.g. as antagonists or to target and/or deliver
CC cytotoxic agents). The polynucleotides are useful therapeutically e.g. to
CC produce antisense sequences to inhibit polypeptide production. They can
CC be used to produce probes and primers useful to detect or isolate
CC sequences encoding PRO polypeptides or similar sequences e.g. variants or
CC sequences from other species. They are also useful for gene mapping and
CC to generate transgenic animals. ABG91355-ABG91363 represent human PRO
CC amino acid sequences of the invention.
XX
XX Sequence 310 AA;
SQ
Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7, 2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFFLLFRGLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKIODEQTYTFDNKIOGLAGRAEILGKTSLKINWTRDSALYRCVVAR 120
DB 61 SDPRIWKIODEQTYTFDNKIOGLAGRAEILGKTSLKINWTRDSALYRCVVAR 120
QY 121 NDRKEIDEIVIELTVQKVPVPCVRPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVIELTVQKVPVPCVRPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRNSSFHNLSETGLVFTAVHKDSDGGYYCIASNDAGSARCEQMEYVDL 240
DB 181 PTDSRANPRNSSFHNLSETGLVFTAVHKDSDGGYYCIASNDAGSARCEQMEYVDL 240
QY 241 NIGGIIGGLVLAVALALITLIGICCAVRRGYFINNKODGESYKKNPKGPDGVNYYIRTDDEG 300
DB 241 NIGGIIGGLVLAVALALITLIGICCAVRRGYFINNKODGESYKKNPKGPDGVNYYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310
XX
RESULT 14
ABG92709
ID ABG92709 standard; Protein; 310 AA.
XX
AC ABG92709;
XX
DT 18-NOV-2002 (first entry)
XX
DE Human secreted protein PRO1868.
XX
KW Human; secreted and transmembrane protein; PRO1800; PRO539;
KW PRO982; PRO1434; PRO1863; PRO1917; PRO1868; PRO3434; PRO1927;
KW inflammatory disorder; immune related disease; rheumatoid arthritis;
KW systemic lupus erythematosus; systemic sclerosis; thyroiditis;
KW autoimmune haemolytic anaemia; diabetes mellitus; infectious hepatitis;
KW psoriasis; allergic disease of the lung; graft-versus host disease;
KW tumour; gene therapy.
XX
OS Homo sapiens.
XX
PN US2002098506-A1.
XX
PD 25-JUL-2002.
XX
PF 27-DEC-2001; 2001US-0033301.
XX
PR 04-AUG-1998; 98US-095325P.
PR 16-DEC-1998; 98US-112851P.
PR 16-DEC-1998; 98US-113145P.
PR 22-DEC-1998; 98US-113511P.
PR 12-JAN-1999; 99US-115558P.
PR 12-JAN-1999; 99US-115565P.
PR 09-FEB-1999; 99US-119341P.
PR 10-FEB-1999; 99US-119537P.
PR 12-FEB-1999; 99US-119565P.
PR 29-OCT-1999; 99US-162506P.
PR 02-JUN-1999; 99WO-US12252.
PR 01-DEC-1999; 99WO-US28634.
PR 02-DEC-1999; 99WO-US28551.
PR 11-FEB-2000; 2000WO-US03565.
PR 22-FEB-2000; 2000WO-US04414.
PR 30-MAR-2000; 2000WO-US05841.
PR 30-MAR-2000; 2000WO-US08439.
PR 30-MAY-2000; 2000WO-US14941.
PR 02-JUN-2000; 2000WO-US15264.
PR 01-DEC-2000; 2000WO-US32678.
XX
XX (GETH ) GENENTECH INC.
XX
XX Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
PI Wood WI;
XX
XX WPI; 2002-690475/74.
DR N-PSDB; ABS68392.
XX
XX Novel secreted and transmembrane polypeptides and polynucleotides
```

PT useful for diagnosis and treatment of inflammatory disorders and
FI immune-related diseases, and identifying modulators -

Claim 12; Fig 14; 125pp; English.

XX The invention relates to an isolated polypeptide having at least 80%
CC amino acid sequence identity to secreted and transmembrane polypeptides
CC PRO1800, PRO539, PRO982, PRO1434, PRO1863, PRO1868, PRO3434 or
CC PRO1927 and their encoding nucleic acids. Also included are vectors, host
CC cells and antibodies against PRO polypeptides. PRO proteins are useful
CC for identifying modulators of the polypeptide. PRO1868 useful for the
CC diagnosis and treatment of inflammatory and immune related diseases
CC including systemic lupus erythematosus, rheumatoid arthritis, systemic
CC sclerosis, autoimmune haemolytic anaemia, thyroiditis, diabetes mellitus,
CC infectious hepatitis, psoriasis, allergic diseases of the lung and
CC graft-versus host disease and tumours. PRO nucleic acids are useful for
CC constructing hybridisation probes for mapping the gene that encodes that
CC PRO and for the genetic analysis of individuals with genetic disorders,
CC and for generating transgenic animals which are useful in the development
CC and screening of therapeutically useful reagents. PRO nucleic acids are
CC also useful for gene therapy, chromosome identification, and tissue
CC typing. PRO proteins are useful as molecular weight markers for protein
CC electrophoresis purposes. The anti-PRO antibodies are useful in
CC diagnostic assays for PRO, e.g. detecting its expression in specific
CC cells, tissues or serum and for affinity purification of PRO.
XX The present sequence represents a PRO protein.

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
DB 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQMEVYDL 240
QY 241 NIGGIIGGLVVLAVLALITLIGICCAVRRGYFINNKQDGESYKXNPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGGLVVLAVLALITLIGICCAVRRGYFINNKQDGESYKXNPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

RESULT 15

ID ABG65296
XX ABG65296 standard; Protein; 310 AA.

XX AC ABG65296;

XX XX 27-AUG-2002 (first entry)

XX DE Human albumin fusion protein #1971.

XX KW Albumin fusion protein; therapeutic protein X; human albumin; HA;
KW human serum albumin; HSA; cancer; reproductive disorder;
KW digestive disorder; immune disorder; endocrine disorder;
KW haematopoietic disorder; neural disorder; connective disorder;
KW cytostatic; antiinfectility; antiinflammatory; antiulcer;

KW immunomodulator; anti-HIV; antidiabetic; haemostatic; nootropic;
KW neuroprotective; antiparkinsonian; antimicrobial; neuroleptic;
KW osteopathic; antiarthritic.

OS Homo sapiens.
OS Synthetic.

XX WO200177137-A1.

XX 18-OCT-2001.

XX 12-APR-2001; 2001WO-US11988.

XX 12-APR-2000; 2000US-229358P.

XX 25-APR-2000; 2000US-199384P.

XX 21-DEC-2000; 2000US-256931P.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Rosen CA, Haseltine WA;

XX WPI; 2002-010886/01.

XX New fusion protein for treating disease e.g. diabetes comprises an

XX PT albumin fused to a therapeutic protein -

XX Claim 1; Page 1893-1894; 2102pp; English.

XX The present invention relates to albumin fusion proteins comprising a
CC therapeutic protein X and human albumin (HA), also known as human serum
CC albumin, HSA). The proteins are useful for treating a disease or
CC disorder that may be modulated by therapeutic protein X. The albumin
CC extends the shelf-life of protein X, and may increase its biological
CC in vitro/in vivo activity. The protein is useful for treating and
CC diagnosing disorders such as cancer, reproductive disorders, digestive
CC disorders (e.g. Crohn's disease, ulcerative colitis), immune disorders
CC (e.g. acquired immunodeficiency syndrome, AIDS), endocrine disorders
CC (e.g. diabetes), haematopoietic disorders, neural disorders
CC (e.g. Alzheimer's, Parkinson's, Creutzfeldt-Jacob disease,
CC encephalomyelitis, meningitis, schizophrenia), and connective disorders
CC (e.g. osteoporosis, arthritis). ABG63326-ABG65518 represent albumin
CC fusion proteins of the invention.

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
DB 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQESGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYYCIASNDAGSARCEQMEVYDL 240
QY 241 NIGGIIGGLVVLAVLALITLIGICCAVRRGYFINNKQDGESYKXNPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGGLVVLAVLALITLIGICCAVRRGYFINNKQDGESYKXNPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

Search completed: December 15, 2003, 14:51:09
Job time : 43 secs
